LIFEPAK® 20e Defibrillator/Monitor CONFIRMATION SHEET



By signing below and returning to Physio-Control, you have acknowledged that you have received the notification letter titled "URGENT MEDICAL DEVICE CORRECTION – ACTION REQUIRED, LIFEPAK® 20e Defibrillator/Monitor" and that it has been delivered to sites, trainers and users of the LIFEPAK 20e device at your facility.

Account #: {End User} {Name} {City, State, Zip} Attention: Risk Management		Completed By (Print Name): Signature: Phone #: (Please return completed form: • By fax to: +31 43 808 0003 • By email to: rsEMEAFA278@stryker.com • Or by mail to: Physio-Control Operations Netherlands B.V. Galjoenweg 68 6222 NV Maastricht The Netherlands		
Serial Number	Confirme Possessi		Device permanently disposed (scrapped) or retired from use	Device cannot be located	Device transferred to another location*		*Please provide the new address and new contact information	
{EXAMPLE}								

Page 1 of 1 February 2018